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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
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NEWS	3	SEP 01	New pricing for the Save Answers for SciFinder Wizard within STN Express with Discover!
NEWS	4	OCT 28	KOREAPAT now available on STN
NEWS	5	NOV 30	PHAR reloaded with additional data
NEWS	6	DEC 01	LISA now available on STN
NEWS	7	DEC 09	12 databases to be removed from STN on December 31, 2004
NEWS	8	DEC 15	MEDLINE update schedule for December 2004
NEWS	9	DEC 17	ELCOM reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	10	DEC 17	COMPUAB reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	11	DEC 17	SOLIDSTATE reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	12	DEC 17	CERAB reloaded; updating to resume; current-awareness alerts (SDIs) affected
NEWS	13	DEC 17	THREE NEW FIELDS ADDED TO IFIPAT/IFIUDB/IFICDB
NEWS	14	DEC 30	EPFULL: New patent full text database to be available on STN
NEWS	15	DEC 30	CAPLUS - PATENT COVERAGE EXPANDED
NEWS	16	JAN 03	No connect-hour charges in EPFULL during January and February 2005
NEWS	17	FEB 25	CA/CAPLUS - Russian Agency for Patents and Trademarks (ROSPATENT) added to list of core patent offices covered
NEWS	18	FEB 10	STN Patent Forums to be held in March 2005
NEWS	19	FEB 16	STN User Update to be held in conjunction with the 229th ACS National Meeting on March 13, 2005
NEWS	20	FEB 28	PATDPAFULL - New display fields provide for legal status data from INPADOC
NEWS	21	FEB 28	BABS - Current-awareness alerts (SDIs) available
NEWS	22	FEB 28	MEDLINE/LMEDLINE reloaded
NEWS	23	MAR 02	GBFULL: New full-text patent database on STN
NEWS	24	MAR 03	REGISTRY/ZREGISTRY - Sequence annotations enhanced
NEWS	25	MAR 03	MEDLINE file segment of TOXCENTER reloaded
NEWS EXPRESS			JANUARY 10 CURRENT WINDOWS VERSION IS V7.01a, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 10 JANUARY 2005
NEWS HOURS			STN Operating Hours Plus Help Desk Availability
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Enter NEWS followed by the item number or name to see news on that specific topic.

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 14:43:41 ON 09 MAR 2005

=> index bioscience

FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED

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ENTRY	SESSION
0.21	0.21

FULL ESTIMATED COST

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DISSABS, ...' ENTERED AT 14:43:58 ON 09 MAR 2005

75 FILES IN THE FILE LIST IN STNINDEX

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=> sequencing same diazomethane
0* FILE DGENE

=> index bioscience

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2.95	3.16

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=> sequencing and diazomethane

- 1 FILE AGRICOLA
- 1 FILE ANABSTR
- 2 FILE BIOSIS
- 1 FILE BIOTECHNO
- 1 FILE CABA
- 7 FILE CAPLUS
- 1 FILE CEN

23 FILES SEARCHED...

- 1 FILE DISSABS
- 1 FILE EMBASE

36 FILES SEARCHED...

- 2 FILE IFIPAT
- 4 FILE MEDLINE
- 1 FILE PASCAL

55 FILES SEARCHED...

- 1 FILE PROMT

2 FILE SCISEARCH
1 FILE TOXCENTER
203 FILE USPATFULL
22 FILE USPAT2
1 FILE WATER
72 FILES SEARCHED...

18 FILES HAVE ONE OR MORE ANSWERS, 75 FILES SEARCHED IN STNINDEX

L1 QUE SEQUENCING AND DIAZOMETHANE

=> d rank

F1	203	USPATFULL
F2	22	USPAT2
F3	7	CAPLUS
F4	4	MEDLINE
F5	2	BIOSIS
F6	2	IFIPAT
F7	2	SCISEARCH
F8	1	AGRICOLA
F9	1	ANABSTR
F10	1	BIOTECHNO
F11	1	CABA
F12	1	CEN
F13	1	DISSABS
F14	1	EMBASE
F15	1	PASCAL
F16	1	PROMT
F17	1	TOXCENTER
F18	1	WATER

=> file caplus medline biosis scisearch
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ENTRY	SESSION
2.36	5.52

FULL ESTIMATED COST

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=> sequencing and diazomethane

L2 15 SEQUENCING AND DIAZOMETHANE

=> dup remove

ENTER L# LIST OR (END):l2

PROCESSING COMPLETED FOR L2

L3 8 DUP REMOVE L2 (7 DUPLICATES REMOVED)

=> d ti 1-8

L3 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1
TI Development of an analytical scheme for simazine and 2,4-D in soil and
water runoff from ornamental plant nursery plots

L3 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN
 TI Purification and partial amino acid sequences of an esterase from tomato

L3 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 2
 TI Cathepsin B-like cysteine proteases confer intestinal cysteine protease activity in *Haemonchus contortus*

L3 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN
 TI Covalent modification of 2'-hydroxyl groups of RNA

L3 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3
 TI Processing the procarboxypeptidase A and other zymogens in murine mast cells

L3 ANSWER 6 OF 8 MEDLINE on STN
 TI The catecholamine binding site of the beta-adrenergic receptor is formed by juxtaposed membrane-spanning domains.

L3 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN
 TI Specific termination of RNA polymerase synthesis as a method of RNA and DNA **sequencing**

L3 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN
 TI Peptide **sequencing** by low-resolution mass spectrometry. I. Use of Acetylacetyl derivatives to identify N-terminal residues

=> d ab bib 2, 8

L3 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN
 AB Screening of 18 suspension plant cell cultures of taxonomically distant species revealed that a Me jasmonate hydrolyzing enzyme activity (0.21-5.67 pkat/mg) occurs in all species so far analyzed. The Me jasmonate hydrolyzing esterase was purified from cell cultures of *Lycopersicon esculentum* using a five-step procedure including anion-exchange chromatog., gel-filtration and chromatog. on hydroxylapatite. The esterase was purified 767-fold to give an almost homogeneous protein in a yield of 2.2%. The native enzyme exhibited a Mr of 26 kDa (gel-filtration chromatog.), which was similar to the Mr determined by SDS-PAGE and MALDI-TOF anal. (Mr of 28547 kDa). Enzyme kinetics revealed a Km value of 15 µM and a Vmax value of 7.97 nkat/mg, an pH optimum of 9.0 and a temperature optimum of 40 °C. The enzyme also efficiently hydrolyzed Me esters of abscisic acid, indole-3-acetic acid, and fatty acids. In contrast, Me esters of salicylic acid, benzoic acid and cinnamic acid were only poor substrates for the enzyme. N-Methylmaleimide, iodoacetamide, bestatin and pepstatin (inhibitors of thiol-, metal- and carboxyproteases, resp.) did not inactivate the enzyme while a serine protease inhibitor, phenylmethylsulfonyl fluoride, at a concentration of 5 mM led to irreversible and complete inhibition of enzyme activity. Proteolysis of the pure enzyme with endoproteinase LysC revealed three peptide fragments with 11-14 amino acids. N-Terminal **sequencing** yielded an addnl. peptide fragment with 10 amino acids. Sequence alignment of these fragments showed high homologies to certain plant esterases and hydroxynitrile lyases that belong to the α/β hydrolase fold protein superfamily.

AN 2002:382784 CAPLUS
 DN 138:85391
 TI Purification and partial amino acid sequences of an esterase from tomato
 AU Stuhlfelder, Christiane; Lottspeich, Friedrich; Mueller, Martin J.
 CS Julius-von-Sachs-Institute for Biosciences, Pharmaceutical Biology, University of Wurzburg, Wurzburg, D-97082, Germany
 SO Phytochemistry (2002), 60(3), 233-240

CODEN: PYTCAS; ISSN: 0031-9422

PB Elsevier Science Ltd.

DT Journal

LA English

RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2005 ACS on STN

AB A description was given of a low-resolution, mass-spectrometric method for the **sequencing** of acetylacetyl (ACA) peptides (I) which gave reliable results with small (2-10 amino acid residues) I, regardless of the amino acids present, and depending on the identification, in the mass spectra visualized, of the N-terminal amino acid residue, "A," which has been found to represent the most prominent peak in the spectra of ACA I esters with N-terminal aliphatic or acidic amino acids. The prominence of these "A" ions in the system employed afforded an unambiguous starting point in the search for the sequence ions (B, C, D, and B1, C1, D1). Under the exptl. conditions employed, arginine I were converted to δ -N-(2-pyrimidinyl) ornithine I, and the ϵ -amino group of lysine was also derivatized; all other functional groups present in the protein amino acids remained intact and were left unprotected. In some cases, partial loss of the side chain was observed with N-terminal methionine, serine, threonine, aspartic acid, and glutamic acid. Some larger seryl- and threonyl-I tended to dehydrate at higher probe temps., making it difficult to recognize these residues in the N-terminal position. Histidyl-, tyrosyl-, phenylalanyl-, and tryptophyl-I showed some elimination of the side chain as ArCH_2^+ or ArCH_2O^+ , but these ions helped to confirm the presence of these residues. Lysyl- and arginyl-I yielded very characteristic [A-99] fragments, and showed only very small "A" fragments in the mass spectra, while cystine derivs. underwent SS bond rupture, accompanied by H^+ transfer. Although I containing unmodified asparagine have been sequenced, **diazomethane** reportedly has to be used instead of alc. HCl for esterification, to prevent hydrolysis to the corresponding aspartyl-I. The ACA-I esters have a relatively high vapor pressure and yielded readily interpretable mass spectra from hepta- and octa-I containing only the neutral amino acids. The presence of basic, polyfunctional amino acids decreased the volatility and limited the sequence procedure to tetra- and penta-I. Expts. to increase the volatility of the ACA-I esters by permethylation reportedly invariably yielded a complex mixture of products, the potential difficulty being the enamino ketone function, which reacts with MeI to give O and C alkylation and a nonvolatile quaternary salt.

AN 1970:51653 CAPLUS

DN 72:51653

TI Peptide **sequencing** by low-resolution mass spectrometry. I. Use of Acetylacetyl derivatives to identify N-terminal residues

AU Bacon, V.; Jellum, E.; Patton, W.; Pereira, W.; Halpern, B.

CS Med. Center, Stanford Univ., Stanford, CA, USA

SO Biochemical and Biophysical Research Communications (1969), 37(6), 878-82

CODEN: BBRCA9; ISSN: 0006-291X

DT Journal

LA English